

1 We claim:

1 1. A method of characterizing a biological specimen, comprising:

2 a) grouping a very large first plurality of entities into a second plurality of groups, each group
3 comprising a small number of entities;

4 b) characterizing each group of entities in the second plurality according to an aspect of the
5 vibrational spectrum of each group; and

6 c) statistically analyzing the characteristics of the groups of entities in the second plurality.

1 2. The method of claim 1, wherein the small number is preponderantly one.

1 3. The method of claim 1, wherein the entities are cells.

1 4. The method of claim 1, wherein characterization of each group is the recording of infrared
2 absorption spectra of the entities in each group.

1 5. The method of claim 4, wherein the small number is preponderantly one, and wherein the
2 entities are cells, and wherein the infrared absorption spectrum of each cell is analyzed for
3 indications that the one cell in each group is in a cell division stage.

1 6. The method of claim 5, wherein the results of the statistical analysis is the percentage of the cells
2 in the cell division stage.

1 7. The method of claim 5, wherein the indication that a cell is in a cell division stage is the
2 presence of a signal indicating DNA in the infrared absorption spectra.

1 8. The method of claim 4, wherein the small number is preponderantly one, and wherein entities
2 are grouped according to the fluorescence of the entities in each group.

1 9. A microscope, comprising:

2 infrared optics for imaging infrared light transmitted through a large number of entities on an area
3 of a microscope stage on to a first detector, where the first detector is an infrared array
4 detector; and

5 optics for imaging fluorescence light emitted by the entities on to a second detector, where the
6 second detector is a fluorescence light array detector.

1 10. The microscope of claim 9, further comprising:

2 a first source of infrared light, the infrared light for illuminating the area of the stage;

3 an second source of ultraviolet light, the ultraviolet light for illuminating the area of the stage.

1 11. The microscope of claim 10, wherein:

2 the first detector is an infra-red area array detector for detecting an image of the entities formed
3 by the infrared light transmitted through the entities;

4 and the second detector is an area array detector for detecting an image of the entities formed by
5 the fluorescence light emitted by the entities.

1 12. The microscope of claim 11, wherein the entities are single cells.

1 13. The microscope of claim 12, wherein the infrared absorption spectra of each cell is recorded.

1 14. The microscope of claim 13, wherein the infrared absorption spectrum of each cell is analyzed
2 for indications that the cell is in a cell division stage.

1 15. The microscope of claim 14, wherein the percentage of the cells in the cell division stage is
2 calculated.

1 16. The microscope of claim 14, wherein the indication that a cell is in a cell division stage is the
2 presence of a signal indicating DNA in the infrared absorption spectra.

1 17. An apparatus, comprising:

2 location means for locating a very large number of cells;

3 vibrational spectrum characterization means for characterizing the vibrational spectrum of each of
4 the cells located by the location means.

1 18. The apparatus of claim 17, wherein the vibrational spectrum characterization means comprises
2 a means for generating and for transmitting infrared light through each cell.

1 19. The apparatus of claim 18, wherein the means for generating infrared light comprises a first
2 laser having a first defined infrared wavelength.

1 20. The apparatus of claim 19, wherein the first laser is pulsed when the location means locates a
2 first cell in a position to be characterized by the first laser.

1 21. The apparatus of claim 19, wherein the first defined wavelength comprises a wavelength
2 wherein DNA is highly absorbing.

1 22. The apparatus of claim 21, wherein a second laser having a second infrared wavelength is
2 pulsed to characterize the first cell, wherein the second infrared wavelength comprises a
3 wavelength wherein RNA is highly absorbing

1 23. The apparatus of claim 20, wherein the first defined wavelength comprises a wavelength
2 wherein DNA is highly absorbing.

1 24. The apparatus of claim 18, wherein the means for generating infrared light comprises a third
2 laser having a broad band infrared wavelength range.

1 25. The apparatus of claim 24, wherein the third laser is pulsed when the location means locates
2 a first cell in a position to be characterized by the laser.

1 26. The apparatus of claim 25, wherein the broad band infrared wavelength range includes a
2 wavelength wherein DNA is highly absorbing.

1 27. The apparatus of claim 26, wherein the broad band infrared wavelength range includes a
2 wavelength wherein RNA is highly absorbing.

1 28. The apparatus of claim 27, wherein the infrared absorption spectrum of each cell is recorded.

1 29. The apparatus of claim 28, wherein the infrared absorption spectrum of each cell is analyzed
2 for indications that the cell is in a cell division stage.

1 30. The apparatus of claim 29, wherein the percentage of the cells in the cell division stage is
2 calculated.

1 31. The apparatus of claim 30, wherein the indication that a cell is in a cell division stage is the
2 presence of a signal indicating DNA in the infrared absorption spectra.

1 33. The apparatus of claim 17, wherein the location means is a fluorescence activated sorting
2 apparatus

1 34. A method of characterizing a large group of biological cells, comprising:

2 a) separating the cells so that the cells of the large group are preponderantly separated from each
3 other;

4 b) characterizing each cell according to an aspect of the vibrational spectrum each cell; and

5 c) statistically analyzing the characteristics of the groups cells.

1 39. The method of claim 38, wherein the separated cells are located according to the fluorescence
2 of the cells.